## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants:

Peter KUFER, et al.

Title:

LESS IMMUNOGENIC BINDING MOLECULES

Application No.:

10/588,734

Filing Date:

February 16, 2005

Examiner:

Unassigned

Art Unit:

Unassigned

Confirmation No.: Unassigned

## INFORMATION DISCLOSURE STATEMENT UNDER 37 C.F.R. §1.56

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

Submitted herewith on Form PTO/SB/08 is a listing of documents known to Applicants in order to comply with Applicants' duty of disclosure pursuant to 37 C.F.R. §1.56.

A copy of each non-U.S. patent document and each non-patent document is being submitted to comply with the provisions of 37 C.F.R. §1.97 and §1.98.

The submission of any document herewith, which is not a statutory bar, is not intended as an admission that such document constitutes prior art against the claims of the present application or that such document is considered material to patentability as defined in 37 C.F.R. §1.56(b). Applicants do not waive any rights to take any action which would be appropriate to antedate or otherwise remove as a competent reference any document which is determined to be a prima facie art reference against the claims of the present application.

## TIMING OF THE DISCLOSURE

The listed documents are being submitted in compliance with 37 C.F.R. §1.97(b), before the mailing date of the first Office Action on the merits.

## RELEVANCE OF EACH DOCUMENT

Documents B1-B30 were cited in the specification of the present application. Documents B1-B14, B16-B19, and B21-B30 are in English. Document B15 is in German and document B20 is in Japanese. Relevance for documents B15 and B20 can be found in the specification of the present application.

No English translation is available for documents B15 or B20; however English abstracts are provided. The absence of a translation does not relieve the PTO from its duty to consider the submitted document (37 CFR §1.98 and M.P.E.P. § 609).

Applicants respectfully request that each listed document be considered by the Examiner and be made of record in the present application and that an initialed copy of Form PTO/SB/08 be returned in accordance with MPEP §609.

Although Applicants believe that no fee is required for this Request, the Commissioner is hereby authorized to charge any additional fees which may be required for this Request to Deposit Account No. 19-0741.

Respectfully submitted.

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Attorney for Applicant Registration No. 35,792

PTO/SB/08 (09-06)

Approved for use through 03/31/2007, OMB 0651-0031

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number.								
		Substitute for form	1449/	PTO	Complete if Known			
	IN.	FORMATION DIS	SCLO	SURE	Application Number	10/588,734		
	s	TATEMENT BY A	PPLI	CANT	Filing Date	2/16/2005		
	_	-to Cubacittadi An	، مانہ	2007	First Named Inventor	Peter KUFER		
Date Submitted: April 6, 2007					Art Unit	Unassigned		
	(use	as many sheets	as ne	cessary)	Examiner Name	Unassigned		
Sheet		1	of	2	Attorney Docket Number	028622-0155		
SHEET			01	, 2	Attorney Bocket Humber	020022-0100		

U.S. PATENT DOCUMENTS						
Examiner	Cite	Document Number	Publication Date	Name of Patentee or Applicant of	Pages, Columns, Lines, Where Relevant	
Initials*	No.1	Number-Kind Code <sup>2</sup> (if known)	MM-DD-YYYY	Cited Document	Passages or Relevant Figures Appear	
	B1	5,565,322 A	10-15-1996	Heller		
	B2	5.585.097 A	12-17-1996	Bolt et al.		
	В3	5,834,597 A	11-10-1998	Tso et al.		
	B4	5,859,205 A	01-12-1999	Adair et al.		
	B5	5,885,573 A	03-23-1999	Bluestone et al.		
	B6	5,929,212 A	07-27-1999	Jolliffe et al.		
	B7	6.407.213 B1	06-18-2002	Carter et al.		
	B8	2002/0131968 A1	09-19-2002	Waldmann et al.		
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			FOREIGN PAT	ENT DOCUMENTS		
Examiner Initials*	Cite No.1	Foreign Patent Document Country Code <sup>3</sup> Number <sup>4</sup> Kind Code <sup>5</sup> (# known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Documents	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T⁰
	B9	EP 0 460 167 B1	10-11-1995	Celltech Therapeutics Limited		
-	B10	EP 0 519 596 B1	02-23-2005	Merck & Co., Inc. National Institutes of Health		
	B11	EP 0 592 106 A1	04-13-1994	Immunogen, Inc.		1
	B12	EP 0 626 390 B1	11-14-2001	Celltech Therapeutics Limited		
	B13	EP 0 940 468 A1	09-08-1999	Genentech, Inc.		
	B14	EP 0 971 959 B1	12-28-2005	Genentech, Inc.		
	B15	EP 1,025,854	08-09-2000	GSF-Forschungszentrum fur Umwelt und Gesundheit GmbH		
	B16	WO 91/09967 A1	07-11-1991	Celltech Limited		
	B17	WO 91/09968 A1	07-11-1991	Celltech Limited		
	B18	WO 92/22653 A1	12-23-1992	Genentech, Inc.		
	B19	WO 00/05268 A1	02-03-2000	BTG International Ltd.		
	B20	WO 03/04648 A1	01-16-2003	Institute of Genetics and Devel. Biol. (Beijing ABT Genetic Engineering Technology Co., Ltd.)		

Examiner Signature		Date Considered	
*EXAMINER: Initial	if reference considered, whether or not citation is in conformance with MPEP 609. Dra-	w line through citation if not in	conformance and not considered.

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•	Substitute for form	1449/	то	Complete If Known		
	INFORMATION DI	SCLO	SURE	Application Number	10/588,734	
	STATEMENT BY	APPLI	CANT	Filing Date	2/16/2005	
	Date Submitted: A	oril 6	2007	First Named Inventor	Peter KUFER	
	Date Submitted. A	DIII O.	2001	Art Unit	Unassigned	
	(use as many sheets	as ne	cessary)	Examiner Name	Unassigned	
Sheet	2	of	2	Attorney Docket Number	028622-0155	

		NON PATENT LITERATURE DOCUMENTS	
Examiner Initials*	Cite No.1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.) date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>6</sup>
	B21	ADAIR et al., "Humanization of the Murine Anti-Human CD3 Monocional Antibody OKT3," Hum. Antibod. Hybridomas, vol. 5, 1 and 2, 1994, pp. 41-47.	
	B22	BRUHL et al., "Depletion of CCR5-Expressing Cells with Bispecific Antithodies and Chemokine Toxins: a New Strategy in the Treatment of Chronic Inflammatory Diseases and HIV," <i>Journal of Immunology</i> , vol. 166, no. 4, Feb. 15, 2001, pp. 2420-2426.	
	B23	HOFFMAN et al., "Lysis of Anti-T3-Bearing Murine Hybridoma Cells by Human Allospecific Cytotoxic T Cell Clones and Inhibition of that Lysis by Anti-T3 and Anti-LFA-1 Antibodies," <i>The Journal of Immunology</i> , vol. 135, no. 1, Jul. 1985, pp. 5-8.	
	B24	KIPRIYANOV, et al., "Bispecific CD3 X CD19 Diabody for T Cell-Mediated Lysis of Malignant Human B Cells," Int. J. Cancer, vol. 77, 1998, pp. 763-772.	
	B25	KUFER et al., "Construction and Biological Activity of a Recombinant Bispecific Single-Chain Antibody Designed for Therapy of Minimal Residual Colorectal Cancer," Cancer Immunol Immunother, vol. 45, 1997, pp. 193-197.	
	B26	LÖFFLER, et al. "A Recombinant Bispecific Single-Chain Antibody, CD19 X CD3, Induces Rapid and High Lymphoma-Directed Cytotoxicity by Unstimulated T Lymphocytes," <i>Blood</i> , vol. 95, no. 6, Mar. 15, 2000, pp. 2098-2103.	
	B27	MACK et al., "Biologic Properties of a Bispecific Single-Chain Antibody Directed Against 17-1A (EpCAM) and CD3," The Journal of Immunology, vol. 58, 1997, pp. 3965-3970.	
	B28	RAUM et al., "Anti-Self Antibodies Selected from a Human IgD Heavy Chain Repertoire: a Novel Approach to Generate Therapeutic Human Antibodies Against Tumor-Associated Differentiation Antigens," Cancer Immunol Immunother, vol. 50, 2001, pp. 141-150.	
	B29	RIECHMANN et al., "Reshaping Human Antibodies for Therapy," Nature, vol. 332, Mar. 24, 1988, pp. 323-327.	
	B30	TRAUNECKER et al., "Bispecific Single Chain Molecules (Janusins) Target Cytotoxic Lymphocytes on HIV Infected Cells," The EMBO Journal, vol. 10, no. 12, 1991, pp. 3655-3659.	

Examiner	Date
Signature	Considered

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Texhwiness: initial in retrieval considered, writing in include copy of this form with next communication to applicant. Applicant's unique citation designation number (optional), 2 See Kinds Codes of USPTO Patent Documents at www.uspto.gov or MPEP 901.04. 3 Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3), 4 For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. 5 Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST, 16 if possible, 6 Applicant is to place a check mark here if English language Translation is attached. This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing his burden, should be sent to the Chief Information Officer, U.S. Patient and Trademark Office, P.O. Box 1460, Alexendrie, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patients, P.O. Box 1450. Alexandria, VA 22313-1450.